

K221410 Maestro PlatformDec 2, 2022
200 days to decisionK221410 · Product code: **FQO** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k221410/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Table, Operating-room, Ac-powered (FQO)
Date received	May 16, 2022
Decision date	Dec 2, 2022
Days to decision	200 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Moon Surgical
Location	Paris, FR
Contact	Anne Osdoit
510(k) history	4 submissions · 4 cleared · 2022-2025

REGULATORY CONSULTANT

Consulting firm	Daniel & Daniel Consulting
Contact	Michael Daniel

Regulatory consulting firm that managed this 510(k) submission on behalf of the applicant. Source: [FDA accessdata.fda.gov](https://accessdata.fda.gov)

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